IN THE UNITED STATES DISTRICT COURT FOR THE NORTHERN DISTRICT OF OHIO EASTERN DIVISION

IN RE: NATIONAL PRESCRIPTION OPIATE LITIGATION

THIS DOCUMENT RELATES TO: Track One-B Trial Case No. 1:17-MD-2804

PHARMACY DEFENDANTS' MOTION IN LIMINE NO. 5 TO PRECLUDE EVIDENCE OR ARGUMENT THAT DEFENDANTS COULD HAVE STOPPED DISTRIBUTING SCHEDULE II CONTROLLED SUBSTANCES

The Court should bar Plaintiffs from offering evidence or argument that Defendants' mere act of distributing opioid medications provides a basis for imposing liability. Plaintiffs should not be able to argue that Defendants could have avoided liability by either choosing never to start distributing opioids or by voluntarily ceasing that distribution. The Supreme Court, the Sixth Circuit, and courts across the country have rejected "stop-selling" or "never-start-selling" theories of liability, and this Court should do the same. Plaintiffs should not be allowed to argue or imply to the jury that it can base liability merely on the fact that Defendants distributed opioid medications, even if they did so in substantial compliance with the Controlled Substances Act.

In *Mutual Pharmaceutical Co. v. Bartlett*, 570 U.S. 472 (2013), the Supreme Court considered whether federal law, which "prohibit[ed] generic drug manufacturers from independently changing their drugs' labels," preempted a state law that would have required the generic drug manufacturer to change its drugs' labels (and thus violate federal law). *Id.* at 475.

¹ Indeed, in the Track 1-A litigation, this Court granted Teva's motion *in limine* to preclude argument that certain Defendants "should have stopped selling generic opioids." Evidentiary Order at 50, *In re Nat'l Prescription Opiate Litig.*, 1:17-MD-2804 (N.D. Ohio Jan. 3, 2020)). This motion is based on the same logic, but reaches distribution, not manufacturing, conduct.

The Court concluded such state laws were preempted, and in reaching that conclusion it rejected the argument that the drug manufacturer could have avoided liability by choosing "not to make [the drug] at all." *Id.* at 488. The Court "reject[ed] this 'stop-selling' rationale as incompatible with [its] pre-emption jurisdiction," which presumed that actors are "not required to cease acting altogether in order to avoid liability." *Id.*

In the wake of that precedent, the Sixth Circuit has recognized that the so-called "stop-selling" theory is not a valid theory of liability. See In re Darvocet, Darvon, & Propoxyphene Prods. Liability Litig., 756 F.3d 917, 925 (6th Cir. 2014) ("The Bartlett court rejected the 'stop selling' theory, namely that a generic manufacturer could have avoided the conflict between state and federal law by refraining from selling the drug."); Strayhorn v. Wyeth Pharm., Inc., 737 F.3d 378, 398 (6th Cir. 2013) ("Nor can the plaintiffs proceed on a failure-to-withdraw or stop-selling theory, a theory recently rejected by the Supreme Court in Bartlett."). So too have other federal courts. See, e.g., Drager v. PLIVA USA, Inc., 741 F.3d 470, 477 (4th Cir. 2014) ("[T]he stop selling rationale is an impermissible means of avoiding preemption under Bartlett."); Beswick v. Sun Pharm. Inds., Ltd., 2018 WL 704399, at *7 (W.D.N.Y. Jan. 30, 2018) ("Insofar as Plaintiff's [claims] can be construed as alleging Defendant should have . . . pulled [its drug] products from the market, in Bartlett the Court flatly rejected the so-called 'stop-selling' rationale").

The Sixth Circuit and other federal courts have recognized for similar reasons that a "never-start-selling" theory (*i.e.* a theory of liability resting on the notion that the defendant could have chosen never to start an activity allowed by federal law) is also invalid. *See, e.g., Yates v. Ortho-McNeil-Janssen Pharm., Inc.*, 808 F.3d 281, 300 (6th Cir. 2015) (rejecting argument that "defendants should never have sold" the challenged product "for the same reasons the Supreme Court in *Bartlett* rejected the stop-selling" theory of liability); *Utts v. Bristol-Myers Squibb Co.*,

226 F. Supp. 3d 166, 186 (S.D.N.Y. 2016) (rejecting argument "that the defendants should never

have sold" the at-issue drug); Brazil v. Janssen Research & Dev. LLC, 196 F. Supp. 3d 1351, 1364

(N.D. Ga. 2016) (rejecting argument that "[d]efendants should have never sold [the drug]").

As in Bartlett, federal law expressly allowed Defendants to distribute Schedule II

controlled substances, including the opioid medications at issue in this case, so long as they

substantially complied with the Controlled Substances Act in the process. See, e.g., 21 U.S.C.

§ 822(b). Consequently, based on the logic of *Bartlett* and its progeny, Defendants may not be

held liable based on the mere fact that they chose to exercise their lawful right to distribute the

opioid medications at issue in this case. Plaintiffs should not be allowed to argue that Defendants

could have avoided liability simply by electing not to distribute opioid medications, irrespective

of whether they complied with the Controlled Substances Act while distributing. See, e.g.,

Bartlett, 570 U.S. at 488; Yates, 808 F.3d at 300; Strayhorn, 737 F.3d at 398.

CONCLUSION

For these reasons, Defendants request respectfully that the Court enter an order barring

Plaintiffs from offering evidence or argument that Defendants' mere act of distributing Schedule

II controlled substances—instead of halting distribution or declining to distribute in the first

instance—as a basis for imposing liability.

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Respectfully submitted,

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